

SPIN-SPIN AND SPIN-ROTATION FINE STRUCTURE OF THE METASTABLE  $a^3\Sigma_u^+$  STATES OF MOLECULAR HELIUM

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In a recent series of experiments<sup>a,b</sup>, we have determined term values of all rotational levels of the  $X^+ 2\Sigma_u^+$  ( $\nu^+ = 0$ ) ground vibronic state of  $^4\text{He}_2^+$  with rotational quantum number  $N^+ \leq 19$  at an accuracy of 25 MHz using MQDT-assisted Rydberg-series extrapolation of metastable helium molecules in the  $a^3\Sigma_u^+$  state. The precision of these experiments was limited by the 150 MHz linewidth of the pulsed laser system employed. In order to improve our resolution and possibly observe the spin-rotation splitting in the  $\text{He}_2^+$  ion, we have replaced the pulsed laser by a CW laser system with a bandwidth of 1.5 MHz. This system was used to measure the spin-spin and spin-rotation fine structure of metastable  $\text{He}_2$  in the  $a^3\Sigma_u^+$  ( $\nu'' = 0$ ) state. Metastable helium molecules were produced by striking a discharge in an expansion of neat helium gas. By cooling the source to a temperature of 10 K, the velocity of the molecular beam was reduced to 500 m/s and an experimental Doppler-limited linewidth of 25 MHz was observed. Fine-structure splittings for all rotational levels with  $N'' \leq 27$  have been measured at an accuracy of 5 MHz and, when possible, have been compared to the values reported in earlier investigations.<sup>c,d,e,f</sup> This comparison revealed a discrepancy that increased with increasing values of  $N''$ . To verify our results, we have recently constructed a variation of a classical molecular-beam magnetic-resonance setup that uses a multistage Zeeman decelerator and a RF stripline for de- and repopulation of the  $F_2$  spin-rotational components with  $J'' = N''$ , respectively.

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<sup>b</sup>L. Semeria, P. Jansen, and F. Merkt, *J. Chem. Phys.* **145**, 204301 (2016).

<sup>c</sup>W. Lichten, M. V. McCusker, and T. L. Vierima, *J. Chem. Phys.* **61**, 2200 (1974).

<sup>d</sup>W. Lichten and T. Wik, *J. Chem. Phys.* **69**, 98 (1978).

<sup>e</sup>M. Kristensen and N. Bjerre, *J. Chem. Phys.* **93**, 983 (1990).

<sup>f</sup>I. Hazell, A. Norregaard, and N. Bjerre, *J. Mol. Spectrosc.* **172**, 135 (1995).